Refine Search

Search Results -

Term	Documents
(2 NOT 3).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	26
(L2 NOT L3).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	26

US Pre-Grant Publication Full-Text Database US Patents Full-Text Database US OCR Full-Text Database Database: EPO Abstracts Database JPO Abstracts Database Derwent World Patents Index IBM Technical Disclosure Bulletins

Search:

		Refine Search
Recall Text 👄	Clear	Interrupt

Search History

DATE: Tuesday, November 23, 2004 Printable Copy Create Case

Set Name side by side	Query	Hit Count	Set Name result set
DB=PGPB	R, USPT, USOC, EPAB, JPAB, DWPI, TDBD; THES=ASSIGNEE; PL	UR = YES;	
OP=AND			
<u>L4</u>	L2 not L3	-26	<u>L4</u>
<u>L3</u>	L2 and (primary adj isolate)	7	<u>L3</u>
<u>L2</u>	L1 and (HIV-1)	33	<u>L2</u>
L1	(prime adj boost) adj (vaccine or immunization)	63	L .1

END OF SEARCH HISTORY

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Welcome to DialogClassic Web(tm)
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Logon file001 23nov04 16:10:11
          *** ANNOUNCEMENT ***
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***Beilstein Facts (File 390)
***Beilstein Reactions (File 391)
***F-D-C Gold/Silver Sheet (File 184)
***BIOSIS Toxicology (File 157)
***IPA Toxicology (File 153)
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                   *** RELOADED
***Toxfile (File 156)
REMOVED
***Textile Technology Digest (File 119)
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     >>> of new databases, price changes, etc.
KWIC is set to 50.
HILIGHT set on as ' '
File
      1:ERIC 1966-2004/Jul 21
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     Set Items Description
Cost is in DialUnits
B 155, 5, 73
      23nov04 16:10:31 User259876 Session D693.1
                    0.230 DialUnits File1
            $0.81
     $0.81 Estimated cost File1
     $0.08 INTERNET
     $0.89 Estimated cost this search
     $0.89 Estimated total session cost 0.230 DialUnits
SYSTEM:OS - DIALOG OneSearch
  File 155:MEDLINE(R) 1951-2004/Nov W2
         (c) format only 2004 The Dialog Corp.
 *File 155: Medline will stop updating COMPLETED records on November 17,
2004. Please see HELP NEWS 155 for details.
  File
        5:Biosis Previews(R) 1969-2004/Nov W2
         (c) 2004 BIOSIS
  File 73:EMBASE 1974-2004/Nov W2
         (c) 2004 Elsevier Science B.V.
     Set Items Description
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1 of 27

```
(HIV-1 (W) VACCINE)
          36392 HIV-1
         253917
                 VACCINE
                 (HIV-1 (W) VACCINE)
      S1
              O
S HIV (W) VACCINE
         356490 HIV
         253917 VACCINE
           2077 HIV (W) VACCINE
      S2
S S2 AND (PRIME-BOOST)
           2077 S2
              0 PRIME-BOOST
                 S2 AND (PRIME-BOOST)
      S3
              0
S PRIME-BOOST (W) VACCINE
              0 PRIME-BOOST
         253917 VACCINE
     S4
              O PRIME-BOOST (W) VACCINE
 (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
          32384 PRIME
          12411 BOOST
         253917 VACCINE
          195281 IMMUNIZATION
                 (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
S S2 AND S5
           2077
                 S2
            148
                 S5
              6 S2 AND S5
?
RD
...completed examining records
              2 RD (unique items)
T S7/3, K/ALL
  7/3, K/1
             (Item 1 from file: 155)
DIALOG(R) File 155: MEDLINE(R)
(c) format only 2004 The Dialog Corp. All rts. reserv.
10890917
          PMID: 11024131
 A human immunodeficiency virus prime - boost
                                                 immunization□regimen in□
        induces antibodies that show interclade cross-reactivity and
 neutralize several X4-, R5-, and dualtropic clade B and C primary isolates.
  Verrier F; Burda S; Belshe R; Duliege A M; Excler J L; Klein M;
Zolla-Pazner S
  Veterans Affairs Medical Center and New York University School of
Medicine, New York, New York 10010, USA.
  Journal of virology (UNITED STATES)
                                       Nov 2000,
                                                  74
                                                      (21)
                                                           p10025-33,
               Journal Code: 0113724
ISSN 0022-538X
  Contract/Grant No.: R01-AI 32424; AI; NIAID; R01-AI 36085; AI; NIAID;
R01-HL 59725; HL; NHLBI; +
  Document type: Clinical Trial; Journal Article
  Languages: ENGLISH
  Main Citation Owner: NLM
  Record type: Completed
  A human immunodeficiency virus prime - boost
                                                 immunization□regimen in□
         induces antibodies that show interclade cross-reactivity and
 neutralize several X4-, R5-, and dualtropic clade B and C primary isolates.
```

2 of 27 11/23/04 4:33 PM

A human immunodeficiency virus (HIV) vaccine that will be useful in

diverse geographic regions will need to induce a broad immune response characterized by cross-clade immunity. To test whether a...

... dualtropic viruses (from clade B) and two R5 viruses (from clades B and C). This is the first demonstration of the induction by a candidate HIV vaccine constructed from clade B laboratory strains of HIV of neutralizing activity against R5 and clade C primary isolates. The data suggest that, by virtue of...

```
7/3,K/2 (Item 2 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2004 The Dialog Corp. All rts. reserv.
```

10527392 PMID: 10630788

Human immunodeficiency virus type 1 envelope-specific cytotoxic T lymphocytes response dynamics after prime - boost vaccine regimens with human immunodeficiency virus type 1 canarypox and pseudovirions.

Arp J; Rovinski B; Sambhara S; Tartaglia J; Dekaban G
Robarts Research Institute, London, Ontario, Canada.

Wirel immunology (INITED STATES) 1999 12 (4)

Viral immunology (UNITED STATES) 1999, 12 (4) p281-96, ISSN

0882-8245 Journal Code: 8801552 Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

Human immunodeficiency virus type 1 envelope-specific cytotoxic T lymphocytes response dynamics after prime - boost vaccine regimens with human immunodeficiency virus type 1 canarypox and pseudovirions.

... in Balb/c mice primed with the recombinant canarypox vector, vCP205, encoding HIV-1 gp120 (MN strain) in addition to Gag/Protease (HIB strain). The prime - boost immunization regimens were administered intramuscularly and involved injections of vCP205 followed by boosts with HIV PSV. Previous vaccination strategies solely involving vCP205 had induced good cellular...

... the capability to effectively induce and boost cell-mediated HIV-1-specific responses. In order to observe the immune effects of HIV PSV in a prime - boost immunization strategy, both DHIV vaccineD immunogens required careful titration in vivo. This suggests that careful consideration should be given to the optimization of immunization protocols destined for human use.

```
Set
        Items
                Description
                (HIV-1 (W) VACCINE)
S1
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S2
         2077
                HIV (W) VACCINE
S3
                S2 AND (PRIME-BOOST)
            Ω
S4
            Ω
                PRIME-BOOST (W) VACCINE
S5
          148
                 (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
S6
            6
                S2 AND S5
S7
            2
                RD (unique items)
S S5 AND (HIV-1)
             148
                  S5
           36392
                  HIV-1
                  S5 AND (HIV-1)
      S8
               9
?
RD
...completed examining records
               9 RD (unique items)
S S9 NOT S7
               9 S9
```

3 of 27 11/23/04 4:33 PM

2 S7 S10 7 S9 NOT S7

T S10/3, K/ALL

10/3,K/1 (Item 1 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

16603088 PMID: 15096801

Immunogenicity of HIV-1 Env and Gag in baboons using a DNA prime/protein boost regimen.

Leung Louisa; Srivastava Indresh K; Kan Elaine; Legg Harold; Sun Yide;

Greer Catherine; Montefiori David C; zur Megede Jan; Barnett Susan W

Chiron Corp., Emeryville, California 94608, USA.

AIDS (London, England) (England) Apr 30 2004, 18 (7) p991-1001,

ISSN 0269-9370 Journal Code: 8710219

Document type: Evaluation Studies; Journal Article

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

... the levels of Env-specific lymphoproliferation. CONCLUSIONS: These results highlight the importance of improving the potency of HIV DNA vaccines by enhanced DNA delivery and **prime** - **boost vaccine** technologies to generate more robust immune responses in larger animal models. In addition, care must be taken when immunizations with Env and Gag antigens are...

Descriptors: *AIDS Vaccines--immunology--IM; *Gene Products, env--immunology--IM; *HIV-1 --immunology--IM; *Vaccines, DNA--immunology--IM; Animals; Antibody Affinity; Cell Division--immunology--IM; DNA, Viral --genetics--GE; Gene Products, env--genetics--GE; Gene Products, gag --immunology--IM; HIV Antibodies--biosynthesis--BI; HIV-1 --genetics--GE; Immunization--methods--MT; Immunization, Secondary--methods--MT; Mutagenesis, Insertional; Papio; T-Lymphocytes, Helper-Inducer--immunology--IM

10/3,K/2 (Item 2 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

15078759 PMID: 12928404

Prime-boost vaccination with HIV-1 Gag protein and cytosine phosphate guanosine oligodeoxynucleotide, followed by adenovirus, induces sustained and robust humoral and cellular immune responses.

Tritel Marc; Stoddard Amy M; Flynn Barbara J; Darrah Patricia A; Wu Chang-you; Wille Ulrike; Shah Javeed A; Huang Yue; Xu Ling; Betts Michael R; Nabel Gary J; Seder Robert A

Cellular Immunology Section, National Institute of Allergy and Infectious Diseases, National Institutes of Health, 40 Convent Drive, Bethesda, MD 20892, USA.

Journal of immunology (Baltimore, Md. - 1950) (United States) Sep 1 2003, 171 (5) p2538-47, ISSN 0022-1767 Journal Code: 2985117R

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

... T cell responses, resulting in CD8+ T cell responses far greater in magnitude than Th1 responses. Furthermore, the Th1 and CD8+ T cell responses following prime - boost immunization were seen in both lymphoid and peripheral mucosal organs and were sustained over several months. Together, these data suggest a new immunization approach for elicitation...

Descriptors: *AIDS Vaccines--immunology--IM; *Adjuvants, Immunologic

--administration and dosage--AD; *Gene Products, gag--immunology--IM; *HIV Antibodies--biosynthesis--BI; * **HIV-1** --immunology--IM; *Immunization, Secondary--methods--MT; *Oligodeoxyribonucleotides--immunology--IM

10/3,K/3 (Item 3 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2004 The Dialog Corp. All rts. reserv.

14233468 PMID: 10029244

HIV-1MN recombinant glycoprotein 160 vaccine-induced cellular and humoral immunity boosted by HIV-1MN recombinant glycoprotein 120 vaccine. National Institute of Allergy and Infectious Diseases AIDS Vaccine Evaluation Group.

Gorse G J; Corey L; Patel G B; Mandava M; Hsieh R H; Matthews T J; Walker M C; McElrath M J; Berman P W; Eibl M M; Belshe R B

St. Louis Department of Veterans Affairs, Medical Center, and Saint Louis University, School of Medicine, Missouri 63106, USA.

AIDS research and human retroviruses (UNITED STATES) Jan 20 1999, 15 (2) p115-32, ISSN 0889-2229 Journal Code: 8709376

Contract/Grant No.: N01-AI-05064; AI; NIAID; N01-AI-15106; AI; NIAID; N01-AI-45211; AI; NIAID; +

Document type: Clinical Trial; Controlled Clinical Trial; Journal Article; Randomized Controlled Trial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

We evaluated **prime** - **boost immunization** with two recombinant envelope glycoprotein subunit vaccines (HIV-1MN recombinant gp160 vaccine in alum adjuvant [MN rgp160] and HIV-1MN recombinant gp120 vaccine in alum...

Descriptors: *AIDS Vaccines--immunology--IM; *Acquired Immunodeficiency Syndrome--immunology--IM; *HIV Envelope Protein gp120--immunology--IM; *HIV-1 --immunology--IM; *Vaccines, Synthetic--immunology--IM

10/3,K/4 (Item 4 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

13749043 PMID: 9444999

Induction of neutralizing antibodies to T-cell line-adapted and primary human immunodeficiency virus type 1 isolates with a prime - boost vaccine regimen in chimpanzees.

Zolla-Pazner S; Lubeck M; Xu S; Burda S; Natuk R J; Sinangil F; Steimer K; Gallo R C; Eichberg J W; Matthews T; Robert-Guroff M

Veterans Affairs Medical Center, New York 10010, USA. Zollas01@mcrcr6.med.nyu.edu

Journal of virology (UNITED STATES) Feb 1998, 72 (2) p1052-9, ISSN 0022-538X Journal Code: 0113724

Contract/Grant No.: AI32424; AI; NIAID; AI36085; AI; NIAID

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

Induction of neutralizing antibodies to T-cell line-adapted and primary human immunodeficiency virus type 1 isolates with a prime - boost vaccine regimen in chimpanzees.

Descriptors: *Antibodies, Viral--immunology--IM; *HIV Envelope Protein gp160--immunology--IM; * HIV-1 --immunology--IM; *T-Lymphocytes--immunology--IM; *Viral Vaccines

10/3,K/5 (Item 5 from file: 155)
DIALOG(R)File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

13031342 PMID: 8679292

Safety and immunogenicity of a recombinant HIV type 1 glycoprotein 160 boosted by a V3 synthetic peptide in HIV-negative volunteers.

Salmon-Ceron D; Excler J L; Sicard D; Blanche P; Finkielstzjen L; Gluckman J C; Autran B; Matthews T J; Meignier B; Kieny M P; et al

Hopital Cochin, Paris, France.

AIDS research and human retroviruses (UNITED STATES) Dec 1995, 11 (12) p1479-86, ISSN 0889-2229 Journal Code: 8709376

Document type: Clinical Trial; Clinical Trial, Phase I; Controlled Clinical Trial; Journal Article

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

... first injection. A weak and short-lived envelope-specific CD(4+)-mediated cytotoxic lymphocyte activity was detected at certain time points in few subjects. This **prime** - **boost vaccine** approach using rgp160 followed by a V3 peptide was safe in humans, and was able to elicit high levels of neutralizing antibodies and a strong...

...; standards--ST; Adult; CD4-Positive T-Lymphocytes--immunology--IM; CD8-Positive T-Lymphocytes--immunology--IM; Cytotoxicity, Immunologic; HIV Antibodies--biosynthesis--BI; HIV Envelope Protein gp160; HIV-1 --chemistry--CH; HIV-1 --genetics--GE; Immunization Schedule; Lymphocyte Activation; Lymphocyte Count--methods--MT; Middle Aged; Neutralization Tests--methods--MT; Random Allocation; Recombinant Proteins--immunology--IM

10/3,K/6 (Item 6 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

12070678 PMID: 12396607

Evaluation in rhesus macaques of Tat and rev-targeted immunization as a preventive vaccine against mucosal challenge with SHIV-BX08.

Verrier Bernard; Le Grand Roger; Ataman-Onal Yasemin; Terrat Celine; Guillon Christophe; Durand Pierre-Yves; Hurtrel Bruno; Aubertin Anne-Marie; Sutter Gerd; Erfle Volker; Girard Marc

UMR 2142 CNRS-BioMerieux, ENSL, 69365 Lyon, France. verrier@cervi-lyon.inserm.fr

DNA and cell biology (United States) Sep 2002, 21 (9) p653-8, ISSN 1044-5498 Journal Code: 9004522

Document type: Journal Article

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

... second group of monkey was primed with SFV-tat only and boosted with MVA-tat. A third group received a tat and rev DNA/MVA prime - boost vaccine regimen. Monitoring of anti-Tat and anti-Rev antibody responses or antigen-specific IFN-gamma production, as measured by enzyme-linked immunospot assays revealed no...

; Animals; Genetic Engineering; HIV Infections--prevention and control --PC; **HIV-1** --immunology--IM; Macaca mulatta; SIV--immunology--IM; Semliki forest virus--immunology--IM; Simian Acquired Immunodeficiency Syndrome --prevention and control--PC; Viral Load

10/3,K/7 (Item 7 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

11338461 PMID: 11427275

Immunization of mice with recombinant gp41 in a systemic prime/mucosal

```
induces HIV-1-specific serum IgG and secretory
         protocol
 antibodies.
  Mantis N J; Kozlowski P A; Mielcarz D W; Weissenhorn W; Neutra M R
  GI Cell Biology Laboratory, Enders 1220, Children's Hospital,
Longwood Avenue, 02115, Boston, MA, USA
                                                  (28-29)
                                                             p3990-4001,
  Vaccine (England)
                       Jul
                                    2001, 19
                              16
0264-410X
            Journal Code: 8406899
  Contract/Grant No.: AI-34757; AI; NIAID; AI-35365; AI; NIAID; DK-34854;
DK; NIDDK; F32 AI10009; AI; NIAID; GM39589; GM; NIGMS; HD-17557; HD; NICHD
  Document type: Journal Article
  Languages: ENGLISH
  Main Citation Owner: NLM
  Record type: Completed
  ... inducing (SI) or non-syncytium-inducing (NSI) North American HIV-1
field isolates, but not uninfected cells. Thus, this recombinant antigen
may be useful in prime / boost immunization protocols designed to
induce systemic and mucosal antibodies that recognize multiple primary
HIV-1 isolates.
                          Vaccines--administration
                                                             dosage -- AD; *HIV
  Descriptors:
                  *AIDS
                                                       and
Antibodies--biosynthesis--BI; *HIV Envelope Protein gp41--administration
and dosage--AD; * HIV-1 --immunology--IM; *Immunoglobulin A, Secretory --biosynthesis--BI; *Immunoglobulin G--biosynthesis--BI; AIDS Vaccines
--genetics--GE; Animals; Antibody Specificity; HIV Antibodies--blood--BL;
HIV Envelope Protein gp41--genetics--GE; HIV-1 --genetics--GE; Hemagglutinin Glycoproteins, Influenza Virus--administration and dosage--AD
                       Glycoproteins, Influenza Virus--genetics--GE;
       Hemagglutinin
Hemagglutinin Glycoproteins, Influenza Virus--immunology--IM; Immunity,
Mucosal...
?
                 Description
Set
        Items
                 (HIV-1 (W) VACCINE)
s1
            O
                HIV (W) VACCINE
         2077
S<sub>2</sub>
                S2 AND (PRIME-BOOST)
S3
            Ω
                 PRIME-BOOST (W) VACCINE
S4
            0
          148
                 (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
S5
                S2 AND S5
S6
            6
                RD (unique items)
S7
            2
            9
                S5 AND (HIV-1)
S8
                RD (unique items)
S9
            9
            7
                S9 NOT S7
S10
?
S S5 AND BX08
             148 S5
              33 BX08
               3 S5 AND BX08
     S11
?
...completed examining records
     S12
               1 RD (unique items)
T S12/3, K/ALL
  12/3, K/1
                (Item 1 from file: 155)
DIALOG(R) File 155: MEDLINE(R)
(c) format only 2004 The Dialog Corp. All rts. reserv.
          PMID: 12396607
 Evaluation in rhesus macaques of Tat and rev-targeted immunization as a
 preventive vaccine against mucosal challenge with SHIV- BX08 .
  Verrier Bernard; Le Grand Roger; Ataman-Onal Yasemin; Terrat Celine;
Guillon Christophe; Durand Pierre-Yves; Hurtrel Bruno; Aubertin Anne-Marie;
Sutter Gerd; Erfle Volker; Girard Marc
                    CNRS-BioMerieux,
                                          ENSL,
          2142
                                                              Lyon,
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7 of 27 11/23/04 4:33 PM

```
verrier@cervi-lyon.inserm.fr
  DNA and cell biology (United States) Sep 2002, 21 (9) p653-8, ISSN
            Journal Code: 9004522
1044-5498
  Document type: Journal Article
  Languages: ENGLISH
  Main Citation Owner: NLM
  Record type: Completed
 Evaluation in rhesus macaques of Tat and rev-targeted immunization as a
 preventive vaccine against mucosal challenge with SHIV- BX08 .
  ... second group of monkey was primed with SFV-tat only and boosted with
MVA-tat. A third group received a tat and rev DNA/MVA prime - boost
          regimen. Monitoring of anti-Tat and anti-Rev antibody responses
or antigen-specific IFN-gamma production, as measured by enzyme-linked
immunospot assays revealed no...
... be clearly established. The animals were challenged by the rectal route
9 weeks after the last booster immunization, using 10 MID(50) of a SHIV-
        stock. Postchallenge follow-up of the monkeys included testing
seroconversion to Gag and Env antigens, measuring virus infectivity in PBMC
by cocultivation with noninfected human...
Set
        Items
                Description
                (HIV-1 (W) VACCINE)
S1
               HIV (W) VACCINE
S2
         2077
               S2 AND (PRIME-BOOST)
S3
            0
               PRIME-BOOST (W) VACCINE
S4
            0
          148
                (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
S5
               S2 AND S5
S6
            6
               RD (unique items)
S7
            2
            9
               S5 AND (HIV-1)
S8
            9
               RD (unique items)
S9
            7
               S9 NOT S7
S10
            3
               S5 AND BX08
S11
S12
            1
               RD (unique items)
?
S S5 AND AVIOPOXVIRUS
             148 S5
               0 AVIOPOXVIRUS
               0 S5 AND AVIOPOXVIRUS
     S13
S S5 AND AVIPOXVIRUS
             148 S5
             223 AVIPOXVIRUS
     S14
               3 S5 AND AVIPOXVIRUS
?
...completed examining records
              3 RD (unique items)
T S15/3, K/ALL
               (Item 1 from file: 155)
  15/3, K/1
DIALOG(R) File 155: MEDLINE(R)
(c) format only 2004 The Dialog Corp. All rts. reserv.
11405093
           PMID: 11500430
  Multistage multiantigen heterologous prime
                                                    boost
 Plasmodium knowlesi malaria provides partial protection in rhesus macaques.
  Rogers W O; Baird J K; Kumar A; Tine J A; Weiss W; Aquiar J C; Gowda K;
Gwadz R; Kumar S; Gold M; Hoffman S L
  Malaria Program, Naval Medical Research Center, Silver Spring, Maryland
20910, Bethesda, Maryland 20889, USA. Rogersb@nmrc.navy.mil
  Infection and immunity (United States) Sep 2001, 69 (9) p5565-72,
```

8 of 27

·ISSN 0019-9567 Journal Code: 0246127

Document type: Journal Article

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

Multistage multiantigen heterologous prime boost vaccine for Plasmodium knowlesi malaria provides partial protection in rhesus macaques.

Descriptors: *Antigens, Protozoan--immunology--IM; * Avipoxvirus
--genetics--GE; *Malaria--prevention and control--PC; *Malaria Vaccines;
*Plasmodium knowlesi--immunology--IM; *Vaccines, DNA; Animals; Antibodies,
Protozoan--blood--BL; Antigens, Protozoan--genetics--GE; Antigens,
Protozoan--metabolism--ME; Avipoxvirus --immunology--IM; Immunization,
Secondary--methods--MT; Interferon Type II--biosynthesis--BI; Macaca
mulatta; Malaria Vaccines--administration and dosage--AD; Malaria Vaccines
--immunology--IM; Mice; Parasitemia...

15/3,K/2 (Item 2 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

11220434 PMID: 11257394

Therapeutic vaccines against melanoma and colorectal cancer.

Tartaglia J; Bonnet M C; Berinstein N; Barber B; Klein M; Moingeon P Aventis Pasteur, Research and Development, Willowdale, Canada.

Vaccine (England) Mar 21 2001, 19 (17-19) p2571-5, ISSN 0264-410X

Journal Code: 8406899

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

... of intratumoral administration of recombinant canarypox viruses carrying cytokine genes. Our current focus is on the induction of tumor-specific T-cell responses using a **prime / boost immunization** schedule with a unique vector system derived from the canary pox virus called ALVAC, in which we incorporate genes encoding Tumor Associated Antigens (TAAs) of...

; Antigens, Neoplasm--genetics--GE; Avipoxvirus --genetics--GE; Cancer Vaccines--genetics--GE; Cancer Vaccines--immunology--IM; Clinical Trials; Colorectal Neoplasms--immunology--IM; Cytokines--genetics--GE; Genetic Vectors; Melanoma--immunology--IM; Safety...

15/3,K/3 (Item 3 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

10527392 PMID: 10630788

Human immunodeficiency virus type 1 envelope-specific cytotoxic T lymphocytes response dynamics after prime - boost vaccine regimens with human immunodeficiency virus type 1 canarypox and pseudovirions.

Arp J; Rovinski B; Sambhara S; Tartaglia J; Dekaban G Robarts Research Institute, London, Ontario, Canada.

Viral immunology (UNITED STATES) 1999, 12 (4) p281-96, ISSN

0882-8245 Journal Code: 8801552

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

Human immunodeficiency virus type 1 envelope-specific cytotoxic T lymphocytes response dynamics after prime - boost vaccine[regimens with] human immunodeficiency virus type 1 canarypox and pseudovirions.

```
... in Balb/c mice primed with the recombinant canarypox vector, vCP205,
encoding HIV-1 gp120 (MN strain) in addition to Gag/Protease (HIB strain).
    prime - boost
                           immunization regimens were administered
intramuscularly and involved injections of vCP205 followed by boosts with
HIV PSV. Previous vaccination strategies solely involving vCP205 had
induced good cellular...
           capability to
                            effectively induce and boost cell-mediated
... the
HIV-1-specific responses. In order to observe the immune effects of HIV PSV
in a prime - boost immunization strategy, both HIV vaccine immunogens
required careful titration in vivo. This suggests that careful
consideration should be given to the optimization of immunization protocols
  Descriptors: *AIDS Vaccines--immunology--IM; * Avipoxvirus --immunology
--IM; *HIV Envelope Protein gp120--immunology--IM; *HIV-1--immunology--IM;
*T-Lymphocytes, Cytotoxic--immunology--IM; *Virion--immunology--IM; Animals
; Avipoxvirus --genetics--GE; Cytotoxicity, Immunologic; Gene Products,
gag--genetics--GE; Gene Products, gag--immunology--IM; HIV Envelope
Protein gp120--genetics--GE; HIV Infections--prevention and control...
               Description
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        Items
S1
                (HIV-1 (W) VACCINE)
         2077
               HIV (W) VACCINE
S2
               S2 AND (PRIME-BOOST)
           0
S3
               PRIME-BOOST (W) VACCINE
S4
            0
               (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
S5
         148
               S2 AND S5
S6
           6
            2
               RD (unique items)
S7
               S5 AND (HIV-1)
S8
           9
           9
               RD (unique items)
S9
               S9 NOT S7
           7
S10
           3
               S5 AND BX08
S11
               RD (unique items)
S12
           1
               S5 AND AVIOPOXVIRUS
S13
           0
           3
               S5 AND AVIPOXVIRUS
S14
               RD (unique items)
S15
?
S S5 AND (ATTENUATED (W) VIRAL (W) VECTOR)
            148 S5
          184443 ATTENUATED
          688552 VIRAL
          277793 VECTOR
              2 ATTENUATED (W) VIRAL (W) VECTOR
              O S5 AND (ATTENUATED (W) VIRAL (W) VECTOR)
  (ATTENUATED (W) VIRAL (W) VECTOR)
          184443 ATTENUATED
          688552 VIRAL
          277793 VECTOR
              2 (ATTENUATED (W) VIRAL (W) VECTOR)
?
...completed examining records
              2 RD (unique items)
T S18/3, K/ALL
               (Item 1 from file: 5)
  18/3, K/1
DIALOG(R)File
              5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.
           BIOSIS NO.: 200400337395
0014966606
 Modified immunogenic pneumolysin compositions as vaccines
AUTHOR: Minetti Conceicao (Reprint); Michon Francis; Pullen Jeffrey K;
```

of 27 11/23/04 4:33 PM

```
Polvino-Bodnar Mary Ellen; Liang Shu-Mei; Tai Joseph Y
AUTHOR ADDRESS: Silver Spring, MD, USA**USA
JOURNAL: Official Gazette of the United States Patent and Trademark Office
Patents 1284 (3): July 20, 2004 2004
MEDIUM: e-file
PATENT NUMBER: US 6764686 PATENT DATE GRANTED: July 20, 2004 20040720
PATENT CLASSIFICATION: 424-2361 PATENT ASSIGNEE: Baxter International Inc.
PATENT COUNTRY: USA
ISSN: 0098-1133 _(ISSN print)
DOCUMENT TYPE: Patent
RECORD TYPE: Abstract
LANGUAGE: English
... ABSTRACT: immunity against Streptococcus pneumoniae. The vaccines may be
  compositions in which the modified pneumolysin is conjugated to bacterial
 polysaccharides or may be carried on an attenuated
                                                      viral
                                                               vector . In
  addition, the invention also provides a method of using the non-toxic,
 modified pneumolysin toxoid in order to stimulate antibodies against
  Streptococcus pneumoniae in...
               (Item 2 from file: 5)
  18/3, K/2
DIALOG(R) File
              5:Biosis Previews(R)
(c) 2004 BIOSIS. All rts. reserv.
0012324200
            BIOSIS NO.: 200000042513
Use of a two-phase partitioning system to purify an immunologically
  attenuated
             viral
                     vector
AUTHOR: Mizouni S K (Reprint); Bradley A J (Reprint); Scott M D (Reprint)
AUTHOR ADDRESS: Albany Medical College, Albany, NY, USA**USA
JOURNAL: Blood 94 (10 SUPPL. 1 PART 2): p415b Nov. 15, 1999 1999
MEDIUM: print
CONFERENCE/MEETING: Forty-first Annual Meeting of the American Society of
Hematology New Orleans, Louisiana, USA December 3-7, 1999; 19991203
SPONSOR: The American Society of Hematology
ISSN: 0006-4971
DOCUMENT TYPE: Meeting; Meeting Abstract
RECORD TYPE: Citation
LANGUAGE: English
 Use of a two-phase partitioning system to purify an immunologically
  attenuated viral
                     vector
Set
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                HIV (W) VACCINE
S2
                S2 AND (PRIME-BOOST)
s_3
            0
                PRIME-BOOST (W) VACCINE
S4
            0
               (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
S5
          148
S6
                S2 AND S5
            6
                RD (unique items)
S7
            2
            9
                S5 AND (HIV-1)
S8
            9
                RD (unique items)
S9
            7
                S9 NOT S7
S10
S11
            3
                S5 AND BX08
                RD (unique items)
S12
            1
                S5 AND AVIOPOXVIRUS
S13
            3
                S5 AND AVIPOXVIRUS
S14
            3
                RD (unique items)
S15
                S5 AND (ATTENUATED (W) VIRAL (W) VECTOR)
S16
                (ATTENUATED (W) VIRAL (W) VECTOR)
S17
               RD (unique items)
S18
S S2 AND REVIEW
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of 27 11/23/04 4:33 PM

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2077 S2
         1745373 REVIEW
            227 S2 AND REVIEW
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S S19 NOT PY<2000
Processing
             227 S19
        32704959 PY<2000
             140 S19 NOT PY<2000
RD
...examined 50 records (50)
...examined 50 records (100)
...completed examining records
            97 RD (unique items)
T S21/3,K/1-10
               (Item 1 from file: 155)
  21/3,K/1
DIALOG(R) File 155: MEDLINE(R)
(c) format only 2004 The Dialog Corp. All rts. reserv.
           PMID: 15285711
 Progress towards the use of Listeria monocytogenes as a live bacterial
 vaccine vector for the delivery of HIV antigens.
  Paterson Yvonne; Johnson Ross S
  University of Pennsylvania, 323 Johnson Pavilion, 36th St. and Hamilton
Walk, Philadelphia, PA 19104-6076, USA. yvonne@mail.med.upenn.edu
                                       Aug 2004, 3 (4 Suppl) pS119-34,
  Expert review of vaccines (England)
ISSN 1476-0584 Journal Code: 101155475
  Contract/Grant No.: AI 36657; AI; NIAID
  Document type: Journal Article
  Languages: ENGLISH
  Main Citation Owner: NLM
  Record type: In Process
\dots for the presentation of passenger antigens to the major histocompatability complex class II and class I pathways of antigen
processing and presentation. This article shall review the progress made
in developing this unusual bacterium as a vaccine vector. In mouse models,
recombinant Listeria carrying a number of different antigens have been...
... oral and parenteral immunization, and in the rhesus macaque after oral
immunization indicate that strong cell-mediated immunity can be induced
         these antigens. This review also discusses safety issues
associated with live bacterial vaccine vectors and problems to be overcome
in developing Listeria as a HIV vaccine for human use.
  21/3, K/2
               (Item 2 from file: 155)
DIALOG(R) File 155: MEDLINE(R)
(c) format only 2004 The Dialog Corp. All rts. reserv.
17040682
          PMID: 15285704
 Neutralizing antibody responses to HIV: role in protective immunity and
 challenges for vaccine design.
  Srivastava Indresh K; Ulmer Jeffrey B; Barnett Susan W
  Chiron Vaccines, 4560 Horton Street, Emeryville, CA 94608, USA. indresh
srivastava@chiron.com
  Expert review of vaccines (England)
                                        Aug 2004, 3 (4 Suppl) pS33-52,
                Journal Code: 101155475
ISSN 1476-0584
  Contract/Grant No.: 1-AI-05396; AI; NIAID; 5 PO1 AI48225-03; AI; NIAID;
AI-95367; AI; NIAID
  Document type: Journal Article
  Languages: ENGLISH
```

11/23/04 4:33 PM

Main Citation Owner: NLM Record type: In Process

... a major health problem throughout the world with a high degree of mortality and morbidity. Therefore, there is an urgent need for an effective anti- HIV vaccine. Although the correlates of protective immunity against infection by HIV remain unidentified, recent studies have demonstrated that both humoral and cellular responses are required for... ... of viral replication in the infected host. Finally, cytotoxic T-lymphocytes may facilitate the clearance of virally infected cells. One of the biggest challenges in HIV vaccine development is to design a HIV envelope immunogen that can induce protective neutralizing antibodies effective against the diverse HIV-1 strains that characterize the global pandemic. The focus of this article is to review the importance of antibodies and the strategies that are currently being used for inducing such antibodies.

21/3,K/3 (Item 3 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

16487841 PMID: 15078177

Humoral immunity in HIV-1 exposure: cause or effect of HIV resistance?

Lopalco Lucia

Clinics for Infectious Diseases, San Raffaele Scientific Institute, Milan, Italy. lucia.lopalco@hsr.it

Curr HIV Res (Netherlands) Apr 2004, 2 (2) p127-39, ISSN 1570-162X Journal Code: 101156990

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

... all over the world. Despite the huge effort in basic and applied research, aimed to control virus spread and to design effective therapeutic strategies, an HIV vaccine is not available yet and current therapeutics approaches cannot prevent the infection. To date, both host genetic repertoire, innate and acquired immune responses, viral mutation...

... responses, e.g. the generation of neutralising antibodies directed against common targets, which can play a protective role in virus entry and/or spread. This **review** focuses on naturally occurring humoral responses to HIV exposure/infection. Moreover, whether such antibodies are induced in response to a peculiar scenario of HIV infection...

21/3,K/4 (Item 4 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

16458936 PMID: 15233732

DNA vaccines against human immunodeficiency virus type 1.

Estcourt Marie J; McMichael Andrew J; Hanke Tomas

MRC Human Immunology Unit, Weatherall Institute of Molecular Medicine, Oxford, UK.

Immunological reviews (Denmark) Jun 2004, 199 p144-55, ISSN 0105-2896 Journal Code: 7702118

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: In Process

Development of a vaccine against human immunodeficiency virus type 1 (HIV-1) is the main hope for controlling the acquired immunodeficiency

syndrome pandemic. An ideal **HIV vaccine** should induce neutralizing antibodies, CD4+ helper T cells, and CD8+ cytotoxic T cells. While the induction of broadly neutralizing antibodies remains a highly challenging goal...

... inducing potent cell-mediated responses in animal models, which are now starting to be tested in humans. Naked DNA immunization is one of them. This review focuses on the stimulation of HIV-specific T cells and discusses in the context of the current 'state-of-art' of DNA vaccines, the areas where this technology might assist either alone or as a part of more complex vaccine formulations in the HIV vaccine development.

21/3,K/5 (Item 5 from file: 155)

DIALOG(R) File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

16103737 PMID: 12871196

Bioorganic approaches towards HIV vaccine design.

Wang Lai-Xi

Institute of Human Virology, University of Maryland Biotechnology Institute, University of Maryland, 725 W. Lombard Street, Baltimore, MD 21201, USA. wangx@umbi.umd.edu

Current pharmaceutical design (Netherlands) 2003, 9 (22) p1771-87,

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

Bioorganic approaches towards HIV vaccine design.

... gp120 as a preventive vaccine was not fulfilled. Broadly neutralizing antibodies and HIV-specific cytotoxic T lymphocytes (CTL) are two immune effectors that an effective HIV vaccine may have to elicit. Experiments in animal models have proved that sufficient levels of neutralizing antibodies can clean up the virus and protect the animals from viral challenge. Therefore, the induction of a broadly neutralizing antibody response remains a principal goal in HIV vaccine development. To achieve persistent infection, HIV has evolved elegant strategies to evade host immune surveillance. These include envelope oligomerization, rapid mutation, heavy glycosylation, and conformational...

... to the immune system. This in turn has greatly facilitated a rational design of immunogens capable of eliciting broadly neutralizing antibodies against HIV. The present review provides an overview of the major scientific obstacles we are facing in the development of an effective HIV vaccine, and discusses recent progresses in the field with a focus on current approaches toward a neutralizing antibody-based HIV vaccine. The bioorganic aspects of the approaches are emphasized.

21/3,K/6 (Item 6 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

15633700 PMID: 14765474

HIV vaccines: current challenges and future directions.

Avrett Sam; Collins Chris

Canadian HIV/AIDS policy & law review / Canadian HIV/AIDS Legal Network (Canada) Jul 2002, 7 (1) p1, 20-5, ISSN 1496-399X Journal Code: 101125215

Document type: Newspaper Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

Volume seven of the **Review** will mark the tenth anniversary of the Canadian HIV/AIDS Legal Network with a series of articles that describe past developments and future directions in...

... to develop and provide access to HIV vaccines. It further explains what is required for governments to fulfill their obligations: additional commitment and resources for HIV vaccine development in the context of increased global research and development regarding diseases of the poor; increased support and advocacy for partnerships to develop HIV vaccines; enhanced regulatory capacity in every country to review , approve, and monitor HIV vaccines; and assurance of global supply of, procurement of, delivery of, and access to vaccines in the context of efforts to...

21/3,K/7 (Item 7 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

15482098 PMID: 14598567

Adenoviruses as vectors for HIV vaccines.

Gomez-Roman Victor Raul; Robert-Guroff Marjorie

Basic Research Laboratory, National Cancer Institute, National Institutes of Health, 41 Library Drive, Building 41, Room D804, Bethesda, MD 20892-5055, USA.

AIDS reviews (Spain) Jul-Sep 2003, 5 (3) p178-85, ISSN 1139-6121 Journal Code: 101134876

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

The tropism of adenoviruses (Ad) for mucosal epithelium makes them ideal vectors for the development of recombinant Ad-HIV vaccines. Currently, several Ad-HIV vaccine candidates are being tested in clinical and preclinical trials. Here, we review the progress on the safety, immunogenicity and efficacy of replication-competent and

replication-defective Ad-HIV and Ad-SIV vaccines in animal models, including non...

21/3,K/8 (Item 8 from file: 155)

DIALOG(R) File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

15332580 PMID: 12016434

HIV-1 polymorphism: a challenge for vaccine development - a review

Morgado M G; Guimaraes M L; Galvao-Castro B

Laboratorio de AIDS e Imunologia Molecular, Departamento de Imunologia, Instituto Oswaldo Cruz-Fiocruz, Rio de Janeiro, RJ, 21045-900, Brasil. mmorgado@ioc.fiocruz.br

Memorias do Instituto Oswaldo Cruz (Brazil) Mar 2002, 97 (2) p143-50 ISSN 0074-0276 Journal Code: 7502619

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

HIV-1 polymorphism: a challenge for vaccine development - a review .
...genetic variability and the limited understanding of the immunological

...genetic variability and the limited understanding of the immunological correlates of protection have made this an enormous scientific challenge not overcome so far. In this **review** we presented an updating of HIV-1 subtypes and recombinant viruses circulating in South American countries, focusing mainly on Brazil, as one of the challenges for **HIV** vaccine development. Moreover, we discussed the importance of stimulating

. developing countries to participate in the process of vaccine evaluation, not only testing vaccines according to already...

21/3,K/9 (Item 9 from file: 155)

DIALOG(R) File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

12494788 PMID: 12957818

Lentivirus infections and mechanisms of disease resistance in chimpanzees.

Rutjens Erik; Balla-Jhagjhoorsingh Sunita; Verschoor Ernst; Bogers Willy; Koopman Gerrit; Heeney Jonathan

Department of Virology, the Biomedical Primate Research Centre, Lange Kleiweg 139, 2288 GJ, Rijswijk, The Netherlands.

Frontiers in bioscience - a journal and virtual library (United States) Sep 1 2003, 8 pd1134-45, ISSN 1093-4715 Journal Code: 9709506

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

... late 1980s, as the AIDS epidemic spread globally in humans, the chimpanzee was eagerly looked to for answers concerning effective AIDS therapies and a possible HIV vaccine . Although from the complicated inter-relationship of the AIDS virus with the human immune system, neither an effective vaccine nor a therapy has emerged, one...

... the spotlight has recently been turned once again on to the chimpanzee, in the intense search for the origin of the AIDS epidemic. Here we **review** the history of HIV-1 infection in this species as well as the observations that have led to some of the current leading hypotheses regarding...

21/3,K/10 (Item 10 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

12456227 PMID: 12876900

Forecasting the future of HIV epidemics: the impact of antiretroviral therapies & imperfect vaccines.

Blower S; Schwartz E J; Mills J

AIDS Institute & Department of Biomathematics, David Geffen School of Medicine at UCLA, 10833 Le Conte Avenue, Los Angeles, CA 90095-1766, USA. sblower@mednet.ucla.edu

AIDS reviews (Spain) Apr-Jun 2003, 5 (2) p113-25, ISSN 1139-6121

Journal Code: 101134876

Contract/Grant No.: 5 T32 AI07370; AI; NIAID; RO1 AI41935; AI; NIAID

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

Mathematical models can be used as health policy tools and predictive tools. Here we **review** how mathematical models have been used both to predict the consequences of specific epidemic control strategies and to design epidemic control strategies. We **review** how models have been used to evaluate the potential impact on HIV epidemics of (i) combination antiretroviral therapies (ART) and (ii) imperfect vaccines. In particular

... discuss, in detail, how mathematical models have been used to evaluate the potential impact of prophylactic, live-attenuated and therapeutic HIV vaccines. We show how HIV vaccine models can be used to evaluate the epidemic-level impact of vaccine efficacy, waning in vaccine-induced

```
· immunity, vaccination coverage level, and changes (increases or...
                 Description
 Set
         Items
 s1
             0
                  (HIV-1 (W) VACCINE)
          2077
                 HIV (W) VACCINE
 S2
                 S2 AND (PRIME-BOOST)
 S3
                 PRIME-BOOST (W) VACCINE
 S4
 S5
           148
                  (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
 S6
             6
                 S2 AND S5
 S7
             2
                 RD (unique items)
 S8
             9
                 S5 AND (HIV-1)
 S9
             9
                 RD (unique items)
             7
                 S9 NOT S7
 S10
 S11
             3
                 S5 AND BX08
                 RD (unique items)
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                 S5 AND AVIOPOXVIRUS
 S13
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                 S5 AND AVIPOXVIRUS
             3
 S14
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                 RD (unique items)
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                 S5 AND (ATTENUATED (W) VIRAL (W) VECTOR)
 S16
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                 S2 AND REVIEW
           227
 S19
           140
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 S20
                 RD (unique items)
            97
 S21
 T S21/3, K/11-20
                  (Item 11 from file: 155)
   21/3,K/11
 DIALOG(R) File 155: MEDLINE(R)
 (c) format only 2004 The Dialog Corp. All rts. reserv.
 12385857
            PMID: 12769793
  Immunological approaches for HIV therapy.
   Lori F; Kelly L M; Lisziewicz J
   Research Institute for Genetic and Human Therapy, at IRCCS Policlinico S.
 Matteo, Pavia, Italy. rightpv@tin.it
   Current drug targets. Infectious disorders (Netherlands)
                                                                  Jun 2003, 3
   (2) p171-8, ISSN 1568-0053
                                  Journal Code: 101128002
   Document type: Journal Article; Review; Review, Tutorial
   Languages: ENGLISH
   Main Citation Owner: NLM
   Record type: Completed
    ... of a successful vaccine are the selection of administration route,
 the eventual clinical application. This review will summarize recently
```

heterologous or homologous prime/boost schedules, and the feasibility of developed preventive and therapeutic vaccines, and carefully evaluate the advantages and potential risks for Human Immunodeficiency Virus (HIV) infected patients. Finally, the concept of "autovaccination" will be defined as it represents the basis for the development of our innovative therapeutic antigen presenting cell targeted HIV vaccine . DermaVir is the first topical vaccine, in combination with antiretroviral therapy, to demonstrate immunological and clinical benefits in a relevant animal model (chronically infected rhesus...

```
21/3,K/12
                (Item 12 from file: 155)
DIALOG(R) File 155: MEDLINE(R)
(c) format only 2004 The Dialog Corp. All rts. reserv.
12333531
           PMID: 12699365
HIV vaccines in infants and children: past trials, present plans and
 future perspectives.
 Safrit Jeffrey T
```

Elizabeth Glaser Pediatric AIDS Foundation, Department of Pediatrics, David Geffin School of Medicine, University of California, Los Angeles, USA. jeff@pedaids.org

Current molecular medicine (Netherlands) May 2003, 3 (3) p303-12,

ISSN 1566-5240 Journal Code: 101093076

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

This review will address the recent history in HIV vaccine trials in the pediatric population while giving due respect to the pediatric vaccine successes achieved over the past decades. Success and failure seen when utilizing the neonatal macaque model of SIV infection and the ramifications of these studies will be discussed. The short list of pediatric HIV vaccine trials currently in progress and those in early planning stages will be reviewed. Finally, future perspectives on the impact of a vaccine that could be...

21/3,K/13 (Item 13 from file: 155)

DIALOG(R) File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

12333526 PMID: 12699360

Dendritic cells as a conduit to improve HIV vaccines.

Pope Melissa

Center for Biomedical Research, Population Council, 1230 York Avenue, New York, NY 10021, USA. mpope@popcbr.rockefeller.edu

Current molecular medicine (Netherlands) May 2003, 3 (3) p229-42,

ISSN 1566-5240 Journal Code: 101093076

Contract/Grant No.: AI40877; AI; NIAID; AI52048; AI; NIAID; AI52060; AI; NIAID; HD41752; HD; NICHD

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

Many potential HIV vaccine strategies are being explored in both animal model and human settings. The success of any vaccine relies on relevant antigenic determinants being presented to the...

...vaccine antigen would be greatly enhanced if targeted to the appropriate DCs to ensure optimal presentation to and subsequently activation of the immune system. This **review** will discuss (i) the current status of DC biology, covering distinct DC subsets and stages of activation and how these influence the types of immune responses that are induced, (ii) how DCs can be exploited to improve the efficacy of **HIV** vaccine strategies currently under investigation, (iii) what has been learned from in vivo model systems using DCs, and (iv) future considerations to advance HIV vaccinology.

21/3,K/14 (Item 14 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

12333525 PMID: 12699359

The role of mucosal immunity in prevention of HIV transmission.

Kozlowski Pamela A; Neutra Marian R

GI Cell Biology Research Laboratory, Children's Hospital and Department of Pediatrics, Harvard Medical School, Boston, MA 02115, USA. pamela.kozlowski@TCH.Harvard.edu

Current molecular medicine (Netherlands) May 2003, 3 (3) p217-28,

ISSN 1566-5240 Journal Code: 101093076

Contract/Grant No.: AI34757; AI; NIAID; AI35365; AI; NIAID; AI48133; AI; NIAID

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

... protection against mucosal transmission of HIV or SIV, whereas cytotoxic T cells are required for clearance of mucosal infection and prevention of systemic spread. This review summarizes the roles of IgA and IgG antibodies in preventing mucosal infection by other viral and bacterial pathogens, and then discusses the various mechanisms by...

... locally infected cells through antibody-dependent cell-mediated cytotoxic reactions. The regional nature of mucosal immune responses is reviewed in light of its relevance to HIV vaccine development. We conclude that mucosal immunization should be considered a component of vaccine strategies against HIV.

21/3,K/15 (Item 15 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

12180686 PMID: 12516046

A review of vaccines for HIV prevention.

Mwau Matilu; McMichael Andrew J

MRC Human Immunology Unit, Weatherall Institute of Molecular Medicine, University of Oxford, John Radcliffe Hospital, Oxford OX3 9DS, UK.

journal of gene medicine (England) Jan 2003, 5 (1) p3-10, ISSN 1099-498X Journal Code: 9815764

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

A review of vaccines for HIV prevention.

... attitude and culture that accelerate the spread of HIV/AIDS have had only modest success. There is urgent need for a prophylactic and/or therapeutic HIV vaccine. This is a review of the obstacles and current trends in HIV vaccine development. Copyright 2002 John Wiley & Sons, Ltd.

21/3,K/16 (Item 16 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

12162574 PMID: 11790602

The past, present and future of HIV - vaccine development: a critical view.

Bojak Alexandra; Deml Ludwig; Wagner Ralf

Institute of Medical Microbiology and Hygiene, Franz-Josef-Strauss Allee 11, 93053 Regensburg, Germany.

Drug discovery today (England) Jan 1 2002, 7 (1) p36-46, ISSN 1359-6446 Journal Code: 9604391

Document type: Historical Article; Journal Article; Review; Review, Tutorial

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

The past, present and future of HIV - vaccine development: a critical view.

... to combat AIDS, the global number of HIV-1 infections is still increasing. There is major consent among scientists worldwide, that the

development of successful HIV vaccine strategies requires a profound understanding of the epidemiological principles of a viral pandemic, as well as deep insights into the molecular and immunological mechanisms of HIV pathogenesis. This review provides an overview of past and present developments, as well as future aspects of HIV vaccines, and also provides a summary of current clinical trials...

21/3,K/17 (Item 17 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

12105315 PMID: 12433353

[Present situation regarding development of an HIV vaccine[][] Situacion actual en el desarrollo de una vacuna frente al virus de la inmunodeficiencia humana.

Alcami Jose

Unidad de Inmunopatologia del SIDA. Centro Nacional de Microbiologia. Instituto de Salud Carlos III. Madrid. Espana. ppalcami@isciii.es

Enfermedades infecciosas y microbiologia clinica (Spain) Dec 2002, 20 (10) p511-22, ISSN 0213-005X Journal Code: 9104081

Document type: Journal Article; Review; Review, Tutorial; English Abstract

Languages: SPANISH

Main Citation Owner: NLM Record type: Completed

[Present situation regarding development of an HIV vaccine[][]
The AIDS epidemic continues to advance, and the development of a preventive HIV vaccine has become a major objective for scientific research. An effective vaccine against this virus is not available and complete protection still has not been achieved in animal models. In this review the major challenges related to the development of a vaccine against HIV are analyzed, particularly the mechanisms involved in viral escape from the immune response...

21/3,K/18 (Item 18 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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11907048 PMID: 12103304

HIV/AIDS treatment and HIV vaccines for Africa.

Weidle Paul J; Mastro Timothy D; Grant Alison D; Nkengasong John; Macharia Doris

Division of HIV/AIDS Prevention, National Center for HIV, STD, and TB Prevention, Centers for Disease Control and Prevention, Atlanta, GA 30333, USA. pweidle@cdc.gov

Lancet (England) Jun 29 2002, 359 (9325) p2261-7, ISSN 0140-6736 Journal Code: 2985213R

Comment in Lancet. 2002 Nov 2;360(9343) 1424; Comment in PMID 12424013

Document type: Journal Article; Review; Review, Academic

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

... In resource-poor settings, antiretroviral drugs should be given with use of standardised treatment regimens and streamlined algorithms for monitoring use. A safe and effective HIV vaccine will supplement prevention efforts to protect uninfected people against infection, or might possibly be able to modify the course of HIV infection. Advances have been

... immune response and immunisation to HIV, and new ideas for candidate vaccines have been developed, including several based on HIV-1 strains prevalent in Africa. **HIV** vaccine efficacy trials are needed in Africa

11/23/04 4:33 PM

to determine whether these advances can be translated into clinical and public health benefits. In this **review**, we discuss the prospects for use of treatment and vaccines in resource-poor settings.

21/3,K/19 (Item 19 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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11694444 PMID: 11868934

Vaccine development against HIV-1: current perspectives and future directions.

Edgeworth Rebecca L; San Juan Homero; Rosenzweig Jason A; Nguyen Nang L; Boyer Jean D; Ugen Kenneth E

Department of Medical Microbiology and Immunology, University of South Florida, College of Medicine, Tampa 33612, USA.

Immunologic research (United States) 2002, 25 (1) p53-74, ISSN 0257-277X Journal Code: 8611087

Contract/Grant No.: P01 AI43069; AI; NIAID

Document type: Journal Article; Review; Review, Academic

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

... virus (HIV) is of great urgency, because it is accepted that vaccination is the only means capable of controlling the AIDS pandemic. The foundation of **HIV** vaccine development is the analysis of immune responses during natural infection and the utilization of this knowledge for the development of protective immunization strategies. Initial vaccine

... has focused on the development of appropriate chemical and genetic adjuvants as well as methods of vaccine delivery to improve the host immune response. This **review** summarizes the vaccine strategies that have been tested in both animal models and human clinical trials.

21/3, K/20 (Item 20 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

11689382 PMID: 11863014 Record Identifier: 101297; VF 9.5.6

Ethical issues in HIV vaccine trials in South Africa. Slack C; Lindegger G; Vardas E; Richter L; Strode A; Wassenaar D

School of Psychology, University of Natal, Private Bag X01, Scottsville, 3209 South Africa.

South African journal of science (South Africa) Jun 2000, 96 p291-5, ISSN 0038-2353 Journal Code: 0066654

KIE Bib: AIDS/human experimentation; human experimentation/foreign countries; immunization

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: KIE

Other Citation Owner: KIE; NRCBL

Record type: Completed

Ethical issues in HIV vaccine trials in South Africa.

In this **review** we describe the ethical issues central to local and international debates about **HIV vaccine** trials. These issues include the physiological and psycho-social risks of trial participation, the preventative interventions to be provided to participants, access to treatment for...

Set Items Description

S1 0 (HIV-1 (W) VACCINE)

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2077
                 HIV (W) VACCINE
. S2
                 S2 AND (PRIME-BOOST)
 S3
                 PRIME-BOOST (W) VACCINE
 S4
            0
 S5
           148
                (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
 S6
            6
                 S2 AND S5
 $7
             2
                 RD (unique items)
                 S5 AND (HIV-1)
 S8
             9
             9
 S9
                 RD (unique items)
 S10
             7
                 S9 NOT S7
 S11
             3
                S5 AND BX08
 S12
             1
                RD (unique items)
                S5 AND AVIOPOXVIRUS
 S13
             0
               S5 AND AVIPOXVIRUS
 S14
             3
               RD (unique items)
 S15
             3
            0 S5 AND (ATTENUATED (W) VIRAL (W) VECTOR)
 S16
                (ATTENUATED (W) VIRAL (W) VECTOR)
             2
 S17
           2
                RD (unique items)
 S18
           227
               S2 AND REVIEW
 S19
           140 S19 NOT PY<2000
 S20
 S21
           97
                 RD (unique items)
 ?
 T S21/3, K/21-30
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21/3,K/21 (Item 21 from file: 155)

DIALOG(R) File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

PMID: 11782252 11610104

T-lymphocyte escape during Understanding cytotoxic simian immunodeficiency virus infection.

O'Connor D; Friedrich T; Hughes A; Allen T M; Watkins D

reviews (Denmark) Oct p115-26, ISSN Immunological 2001, 183

Journal Code: 7702118 0105-2896

Document type: Journal Article; Review; Review, Academic

Languages: ENGLISH Main Citation Owner: NLM Record type: Completed

Infection of rhesus macaques with simian immunodeficiency virus (SIV) is excellent model system for studying viral adaptation to immune an responses. In this review , we discuss how the SIV-infected macaque has provided unequivocal evidence for cytotoxic T-lymphocyte (CTL) selection of viral escape variants. This improved understanding of CTL escape may influence human immunodeficiency virus (HIV) vaccine design as well as our understanding of HIV pathogenesis.

```
21/3,K/22
              (Item 22 from file: 155)
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DIALOG(R) File 155:MEDLINE(R)

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PMID: 11708271 11522515

Shooting blanks. "Science" writer Jon Cohen speaks of how the search for an HIV vaccine□ has strayed. Interview by Bob Roehr.□

Cohen J

IAPAC monthly (United States) Sep 2001, 7 (9) p268-70,

Journal Code: 101087241

Document type: Interview; Newspaper Article Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

Shooting blanks. "Science" writer Jon Cohen speaks of how the search for HIVvaccine□ has strayed. Interview by Bob Roehr.□

AIDS Vaccines--standards--ST; Animals; Disease Models, Animal; Drug

2 of 27 11/23/04 4:33 PM Industry--organization and administration--OG; Haplorhini; Health Policy --trends--TD; Journalism, Medical; Leadership; Needs Assessment; Peer Review, Research; United States

21/3,K/23 (Item 23 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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11507946 PMID: 11672892

Rational development of prophylactic HIV vaccines based on structural and regulatory proteins.

Mooij P; Heeney J L

Department of Virology, Biomedical Primate Research Centre, P.O. Box 3306, 2288 Rijswijk, The Netherlands.

Vaccine (England) Nov 12 2001, 20 (3-4) p304-21, ISSN 0264-410X

Journal Code: 8406899

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed

The severity of the AIDS epidemic clearly emphasises the urgent need to expedite HIV vaccine candidates into clinical trials. Prophylactic HIV vaccine candidates have been evaluated in non-human primates. Based on specific proof of principle studies the first phase III clinical studies have recently begun in humans. However, a truly effective HIV vaccine is not yet at hand and many problems related to specific properties of the virus remain to be overcome. Previously proven empirical approaches have largely failed and now rational thinking based on an understanding of immunity to lentiviral infections is needed. This review addresses the scientific problems and complications facing the development of an HIV vaccine as well as the possible strategies currently available to overcome these problems. Recent attention has focussed on identifying the immune correlates and mechanisms of protection...

21/3,K/24 (Item 24 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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11260442 PMID: 11339361

HIV vaccine development at Duke University Medical Center.

Haynes B F; Liao H X; Staats H F; Alam M S; Weinhold K J; Montefiori D C Department of Medicine, The Duke Center for Aids Research, Duke University Medical Center, Durham, NC 27710, USA. hayne002@mc.duke.edu Immunologic research (United States) 2000, 22 (2-3) p263-9, ISSN 0257-277X Journal Code: 8611087

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

HIV vaccine development at Duke University Medical Center.

With the AIDS epidemic continuing to spread throughout the world, development of a safe, practical, and effective HIV vaccine is a national priority. HIV vaccine research efforts are currently targeted towards design of HIV immunogens that induce both cellular and humoral immunity. This brief review summarizes ongoing work at the Duke University School of Medicine on HIV vaccine development.

21/3,K/25 (Item 25 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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10836864 PMID: 10964278

Vaccines for the control of HIV/AIDS.

Gotch F; Rutebemberwa A; Jones G; Imami N; Gilmour J; Kaleebu P; Whitworth J

Department of Immunology, Imperial College of Science and Medicine, London, UK. f.gotch@ic.ac.uk

Tropical medicine & international health - TM & IH (ENGLAND) Jul 2000,

5 (7) pA16-21, ISSN 1360-2276 Journal Code: 9610576 Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

This review discusses the feasibility of an HIV vaccine and describes the history, efficacy and potential to succeed of old and new vaccine concepts.

21/3,K/26 (Item 26 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2004 The Dialog Corp. All rts. reserv.

10833466 PMID: 10963285

Challenges in the development of an effective HIV vaccine : current approaches and future directions.

Klein E; Ho R J

Department of Pharmaceutics, School of Pharmacy, University of Washington, Seattle, 98195-7610, USA.

Clinical therapeutics (UNITED STATES) Mar 2000, 22 (3) p295-314;

discussion 265, ISSN 0149-2918 Journal Code: 7706726

Document type: Journal Article; Review; Review, Tutorial

Languages: ENGLISH

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

Challenges in the development of an effective HIV vaccine : current approaches and future directions.

OBJECTIVE: The intent of this review is to investigate and discuss why developing a successful HIV vaccine has been so challenging, first by examining the molecular biology of the virus and how HIV interacts with the immune system, and then reviewing past viral vaccine successes as well as future directions for HIV vaccine research. BACKGROUND: Since HIV appeared in the United States in the early 1980s, an estimated 40 million people worldwide have been infected with the virus...

... vaccine entering the market in the foreseeable future. METHODS: MEDLINE was searched for articles written between 1966 and June 1999. Search terms used were AIDS, HIV vaccine, HIV-1, HIV-2, vaccines, and human immunodeficiency virus. RESULTS: Only 2 candidates for an HIV vaccine are currently in phase III clinical trials (1 in the United States and 1 in Thailand). The efficacy of these vaccines when applied to the...

21/3,K/27 (Item 27 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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10786278 PMID: 10912619

Listeria monocytogenes as an alternative vaccine vector for HIV.

Mata M; Paterson Y

University of Pennsylvania Medical School, Department of Microbiology, Philadelphia, USA.

Archivum immunologiae et therapiae experimentalis (POLAND) 2000, 48 (3) p151-62, ISSN 0004-069X Journal Code: 0114365

Document type: Journal Article; Review; Review, Academic

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

The necessity for an HIV vaccine and a brief review of current strategies towards this aim are given here to set into context contemporary studies towards exploiting the bacterium Listeria monocytogenes as an HIV vaccine vector. The cell biology and immunology of this unusual intracellular organism are also reviewed, in addition to its application to introducing viral antigens, including HIV...

21/3,K/28 (Item 28 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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10595587 PMID: 10701170 Record Identifier: 64608

Ethical considerations in international HIV vaccine trials: summary of a consultative process conducted by the Joint United Nations Programme on HIV/AIDS (UNAIDS).

Guenter D; Esparza J; Macklin R

Department of Family Medicine, McMaster University, Hamilton, Canada. Journal of medical ethics (ENGLAND) Feb 2000, 26 (1) p37-43, ISSN 0306-6800 Journal Code: 7513619

6 fn.; Full author name: Esparza, Jose; Full author name: Guenter, Dale; Full author name: Macklin, Ruth; KIE BoB Subject Heading: AIDS/human experimentation; KIE BoB Subject Heading: human experimentation/foreign countries

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Other Citation Owner: KIE Record type: Completed

Ethical considerations in international HIV vaccine trials: summary of a consultative process conducted by the Joint United Nations Programme on HIV/AIDS (UNAIDS).

... international consultation; its purpose was further to define the important ethical issues and to formulate guidance that might facilitate the ethical design and conduct of **HIV vaccine** trials in international contexts. This paper summarises the major outcomes of the UNAIDS consultative process.

; AIDS Vaccines; Attitude to Health; Bioethical Issues; Consensus; Control Groups; Ethical Review; Ethics; Ethics, Research; Health Priorities; Moral Obligations; Needs Assessment; Nontherapeutic Human Experimentation; Patient Advocacy; Research Design--standards--ST; Research Subjects; Social Justice; Social Values; Socioeconomic...

21/3,K/29 (Item 1 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)

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0015069003 BIOSIS NO.: 200400436922

HIV vaccine and immunotherapy approaches

AUTHOR: Gruber Andreas (Reprint)

AUTHOR ADDRESS: Neptunusstr 142, NL-2586 GX, Den Haag, Netherlands**

Netherlands

AUTHOR E-MAIL ADDRESS: agruber@ucsd.edu

JOURNAL: Current Medicinal Chemistry - Immunology Endocrine & Metabolic

Agents 4 (1): p21-25 March 2004 2004

MEDIUM: print

ISSN: 1568-0134 _(ISSN print)

DOCUMENT TYPE: Article; Literature Review

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RECORD TYPE: Citation
LANGUAGE: English
       vaccine and immunotherapy approaches
HIV
DESCRIPTORS:
 MISCELLANEOUS TERMS:
                          ...Literature Review
                (Item 2 from file: 5)
  21/3,K/30
DIALOG(R) File
                5:Biosis Previews (R)
(c) 2004 BIOSIS. All rts. reserv.
0015010781
            BIOSIS NO.: 200400381570
 The problems of variability of the human immunodeficiency virus
AUTHOR: Ivans'ka N V; Trokhimchuk T Yu
JOURNAL: Biopolimery i Kletka 20 (3): p171-181 May 2004 2004
MEDIUM: print
ISSN: 0233-7657 (ISSN print)
DOCUMENT TYPE: Article; Literature Review
RECORD TYPE: Abstract
LANGUAGE: Ukrainian
ABSTRACT: This review deals with variability of the human
  immunodeficiency virus (HIV) and the epidemiological consequences of such
  variability. The level of the HIV variability is extremely high...
DESCRIPTORS:
  CHEMICALS & BIOCHEMICALS:
                                HIV
                                      vaccine
?
Set
        Items
                Description
                 (HIV-1 (W) VACCINE)
S1
            0
S2
         2077
                HIV (W) VACCINE
                S2 AND (PRIME-BOOST)
S3
            0
                PRIME-BOOST (W) VACCINE
S4
            0
                (PRIME (W) BOOST) (W) (VACCINE OR IMMUNIZATION)
S5
          148
                S2 AND S5
S6
            6
S7
            2
                RD (unique items)
S8
                S5 AND (HIV-1)
S9
            9
                RD (unique items)
            7
                S9 NOT S7
S10
            3
                S5 AND BX08
S11
S12
            1
                RD (unique items)
                S5 AND AVIOPOXVIRUS
S13
            0
                S5 AND AVIPOXVIRUS
S14
            3
                RD (unique items)
S15
            3
                S5 AND (ATTENUATED (W) VIRAL (W) VECTOR)
S16
            0
S17
            2
                (ATTENUATED (W) VIRAL (W) VECTOR)
S18
            2
                RD (unique items)
                S2 AND REVIEW
S19
          227
                S19 NOT PY<2000
S20
          140
                RD (unique items)
S21
           97
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COST
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            $8.61 41 Types
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                      1.626 DialUnits File5
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            $7.00 4 Types
    $16.11 Estimated cost File5
                     1.153 DialUnits File73
           $11.30
    $11.30 Estimated cost File73
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11/23/04 4:33 PM

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\$5.49 INTERNET \$47.96 Estimated cost this search \$48.85 Estimated total session cost 5.025 DialUnits

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